

on STN

ACCESSION NUMBER: 1999163820 EMBASE
TITLE: Photodynamic diagnosis and treatment for atherosclerosis
by
an endoscopic approach.
AUTHOR: Hayashi J.; Saito T.; Aizawa K.
CORPORATE SOURCE: J. Hayashi, Department Medicine and Gerontology, School of
Medicine, Kyorin University, Shinkawa 6-20-2, Mitakashi,
Tokyo 181, Japan
SOURCE: Diagnostic and Therapeutic Endoscopy, (1999) 5/3
(191-195).

Refs: 9

ISSN: 1070-3608 CODEN: DTENER

COUNTRY: United Kingdom

DOCUMENT TYPE: Journal; Article

FILE SEGMENT: 006 Internal Medicine
014 Radiology
018 Cardiovascular Diseases and Cardiovascular Surgery
027 Biophysics, Bioengineering and Medical
Instrumentation
037 Drug Literature Index

LANGUAGE: English

SUMMARY LANGUAGE: English

AB . . . the carotid artery were recognized as reddish spots from outside
the artery. In addition, we visualized specifically at the beating
heart surface small coronary atherosclerosis using an
epifluorescence stereoscope system. We examined the effects of
photodynamic treatment with NPe6 on the. . .

RN (aspartylchlorin e6) 110230-98-3

AB The photosensitizer, mono-L-aspartyl chlorin e6 (NPe6), specifically
accumulates in the atheromatous plaque. We detected the fluorescence
spectra of NPe6 emitted from atheromatous plaques on the descending
thoracic aorta by an angioscopic approach using the animal model of
atherosclerosis. We also showed that a fluorescence spectrum peak at 675
nm, was obtained laparoscopically only in parts of the abdominal aorta
with an atheromatous plaque. By a fluorescence endoscope, atheromatous
plaques on the carotid artery were recognized as reddish spots from
outside the artery. In addition, we visualized specifically at the
beating

heart surface small coronary atherosclerosis using an
epifluorescence stereoscope system. We examined the effects of
photodynamic treatment with NPe6 on the atheromatous plaque. The change
in

the elastic framework in the atheromatous plaque after photodynamic
treatment was evaluated using scanning electron microscopy. The
destruction of the architecture of the elastic fiber network in the
atheromatous plaque was revealed. We also studied the change in the lipid
components of the atheromatous plaque using Fourier transform infrared
(FTIR) microspectroscopy. FTIR microspectroscopic analysis showed a
dissociation of ester bonds of cholesterol esters in the atheromatous
plaque after photodynamic treatment. The framework of the atheromatous
plaque and the lipids accumulated in the plaque could be destroyed
following such treatment.

FILE 'REGISTRY' ENTERED AT 15:23:55 ON 07 SEP 2004

L1 1 S 110230-98-3/RN

FILE 'CAPLUS, EMBASE, BIOSIS, USPATFULL' ENTERED AT 15:24:21 ON 07 SEP 2004

L2 206 S L1

L3 3 S L2 AND ANGIOPLAS?

L4 6 S L2 AND (HEART OR CARDIO?)

L5 34985 S PCTA OR (PERCUTANEOUS TRANSLUMINAL)

L6 2 S L2 AND L5

L7 6 S L4 NOT L6

L6 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1995:583727 CAPLUS

DOCUMENT NUMBER: 123:78621

TITLE: **Percutaneous transluminal**
photodynamic therapy of atheroma using
mono-L-aspartylchlorin e6

AUTHOR(S): Hayashi, Junichi; Sato, Hideaki; Saito, Takashi;
Kuroiwa, Yukari; Aizawa, Katsuo; Fujiwara, Tatsushi;
Hosoda, Yasuhiro

CORPORATE SOURCE: School Medicine, Kyorin University, Mitakashi, 181,
Japan

SOURCE: Proceedings of SPIE-The International Society for
Optical Engineering (1995), 2371, 554-7
CODEN: PSISDG; ISSN: 0277-786X

DOCUMENT TYPE: Journal

LANGUAGE: English

TI **Percutaneous transluminal** photodynamic therapy of
atheroma using mono-L-aspartylchlorin e6

IT Laser radiation
Photosensitizers

(**percutaneous transluminal** photodynamic therapy of
atheroma using mono-L-aspartylchlorin e6 and laser radiation)

IT Antiarteriosclerotics

(antiatherosclerotics, photosensitizing; **percutaneous**
transluminal photodynamic therapy of atheroma using
mono-L-aspartylchlorin e6 and laser radiation)

IT Phototherapy

(chemo-, **percutaneous transluminal** photodynamic
therapy of atheroma using mono-L-aspartylchlorin e6 and laser
radiation)

IT Photodynamic action

(therapeutic, **percutaneous transluminal**
photodynamic therapy of atheroma using mono-L-aspartylchlorin e6 and
laser radiation)

IT **110230-98-3**, Mono-L-aspartylchlorin e6

RL: BAC (Biological activity or effector, except adverse); BSU

(Biological

study, unclassified); THU (Therapeutic use); BIOL (Biological study);

USES

(Uses)

(**percutaneous transluminal** photodynamic therapy of
atheroma using mono-L-aspartylchlorin e6 and laser radiation)

AB Structural changes after photodynamic therapy of atherosclerotic lesions
of the thoracic aorta were analyzed by SEM. Cholesterol fed
atherosclerotic rabbits were injected i.v. with 5 mg/kg of NPe6. At 6 h
after NPe6 loading, a diode laser irradiated angioscopically on the
surface of atheroma with the total energy of 200 mJ/cm². SEM showed
degeneration of atherosclerotic plaques of the thoracic aorta examd. at
one week after photodynamic therapy. NPe6 could be a potent
photosensitizer for photodynamic therapy of atheroma.

ACCESSION NUMBER: 2000:725466 CAPLUS
 DOCUMENT NUMBER: 133:291111
 TITLE: Inhibitors for vascular recontriction after
 angioplasty
 INVENTOR(S): Nagae, Tsuneyuki; Aizawa, Katsuo
 PATENT ASSIGNEE(S): Meiji Seika, Kaisha, Ltd., Japan
 SOURCE: PCT Int. Appl., 33 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: Japanese
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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WO 2000059505	A1	20001012	WO 2000-JP2156	20000403
W: JP, US				
RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
EP 1166784	A1	20020102	EP 2000-913083	20000403
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI				

PRIORITY APPLN. INFO.:

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